



<b>Office Action Summary</b>	<b>Application No.</b> 10/588,603	<b>Applicant(s)</b> HIRABAYASHI ET AL.
	<b>Examiner</b> STEFANIE COHEN	<b>Art Unit</b> 4162

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --  
**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If no period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED. (35 U.S.C. § 133).

Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

#### **Status**

1) Responsive to communication(s) filed on 09 June 2008.

2a) This action is FINAL.      2b) This action is non-final.

3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

#### **Disposition of Claims**

4) Claim(s) 1,2,5,7 and 8 is/are pending in the application.

4a) Of the above claim(s) \_\_\_\_\_ is/are withdrawn from consideration.

5) Claim(s) \_\_\_\_\_ is/are allowed.

6) Claim(s) 1,2,5,7 and 8 is/are rejected.

7) Claim(s) \_\_\_\_\_ is/are objected to.

8) Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

#### **Application Papers**

9) The specification is objected to by the Examiner.

10) The drawing(s) filed on 07 August 2006 is/are: a) accepted or b) objected to by the Examiner.

Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).

Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).

11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

#### **Priority under 35 U.S.C. § 119**

12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).

a) All    b) Some \* c) None of:

1. Certified copies of the priority documents have been received.
2. Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

#### **Attachment(s)**

1) Notice of References Cited (PTO-892)

2) Notice of Draftsperson's Patent Drawing Review (PTO-948)

3) Information Disclosure Statement(s) (PTO/SB/08)  
Paper No(s)/Mail Date 0/9/2008; 4/17/2008; 8/7/2006.

4) Interview Summary (PTO-413)  
Paper No(s)/Mail Date \_\_\_\_\_.

5) Notice of Informal Patent Application

6) Other: \_\_\_\_\_



Regarding claim 5 and 8, Ogura, ex 13, teach platinum as the catalyst metal.

Claim 2 is rejected under 35 U.S.C. 103(a) as being unpatentable over Ogura et al. (Japanese Unexamined Patent Publication No. 10-258232) in view of Wright (2001). Ogura teaches a catalyst for exhaust gas purification comprising a core part and a catalyst carrying layer. Ogura further teaches the core comprising cerium oxide which is an NO<sub>x</sub> occlusion material and alumina and the catalyst layer comprising platinum supported on alumina and cerium oxide. Ogura, paragraph 25, also teaches it is desirable to have a ratio of .05-10 mol to 1 mol of ceria oxide to alumina a particle size ranging from .1-10 micrometers in the core part. It would have been obvious to one of ordinary skill in the art at the time of the invention to optimize the particle size and the aluminum/ ceria ratio of the catalyst carrying layer to maximize the efficiency of the catalyst. It also would have been obvious to one of ordinary skill in the art at the time of the invention that when the molar number of ceria is greater than the molar number of alumina in the catalyst carrying layer, the alumina is less impeding on the ceria oxygen storage capacities. Although Ogura teaches a composition for a catalyst, Ogura does not teach using sols as starting components to form a catalyst. Wright teaches advantages of using sol- gel synthesized materials. Wright, sect 1.4, teaches advantages of using sol- gel sol- gel synthesized materials including temperatures required for all stages are low and close to room temperature. This results in a minimal thermal degradation and high purity. Wright further teaches control may be achieved

over particle size and pore size, porosity and pore wall chemistry of the final material. It would have been obvious to one of ordinary skill in the art at the time of the invention to use a ceria sol and alumina sol to form the Ogura catalyst because Wright teaches advantages of sols which will result in a highly pure Ogura catalyst.

Claim 7 is rejected under 35 U.S.C. 103(a) as being unpatentable over Ogura et al. (Japanese Unexamined Patent Publication No. 10-258232) in view of Wright (2001) and in further view of Pinnavaia (4981825). Ogura teaches a catalyst for exhaust gas purification comprising a core part and a catalyst carrying layer. Ogura further teaches the core comprising cerium oxide which is an NO<sub>x</sub> occlusion material and alumina and the catalyst layer comprising platinum supported on alumina and cerium oxide. Ogura, paragraph 25, also teaches it is desirable to have a ratio of .05-10 mol to 1 mol of ceria oxide to alumina a particle size ranging from .1-10 micrometers in the core part. It would have been obvious to one of ordinary skill in the art at the time of the invention to optimize the particle size and the aluminum/ ceria ratio of the catalyst carrying layer to maximize the efficiency of the catalyst. It also would have been obvious to one of ordinary skill in the art at the time of the invention that when the molar number of ceria is greater than the molar number of alumina in the catalyst carrying layer, the alumina would be less impeding on the ceria oxygen storage capacities therefore improving the efficiency of the catalyst. Ogura, ex. 13, teaches forming a core part for the catalyst by mixing K<sub>2</sub>O, TiO<sub>2</sub> and Al<sub>2</sub>O<sub>3</sub> and further mixing CeO<sub>2</sub> but does not teach using sols as starting components to form a catalyst. Wright teaches advantages of using sol- gel

synthesized materials. Wright, sect 1.4, teaches advantages of using sol- gel sol- gel synthesized materials including temperatures required for all stages are low and close to room temperature. This results in a minimal thermal degradation and high purity. Wright further teaches control may be achieved over particle size and pore size, porosity and pore wall chemistry of the final material. It would have been obvious to one of ordinary skill in the art at the time of the invention to use a ceria sol and alumina sol to create the Ogura catalyst because Wright teaches advantages of sols which will result in a highly pure Ogura catalyst. Although Ogura teaches a process for producing the catalyst, Ogura does not teach adjusting the pH of the solution. Pinnavaia teaches mixing a silica sol gel useful for catalytic gaseous reactions. Pinnavaia, ex 1, teaches the pH of the silica sol gel was decreased from 10 to 3 by adding HCl. It would have been obvious to one of ordinary skill in the art at the time of the invention to add a calculated amount of HCl to the Ogura core components because this is one specific way of reaching a necessary isoelectric point. It also would have been obvious to one of ordinary skill in the art at the time of the invention that the isoelectric points of ceria and aluminum depend on the starting material of the elements and can easily be adjusted to optimize the performance of the catalyst. Ogura further teaches the solution was stirred, grinded and calcinated to form the core powder part.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to STEFANIE COHEN whose telephone number is

